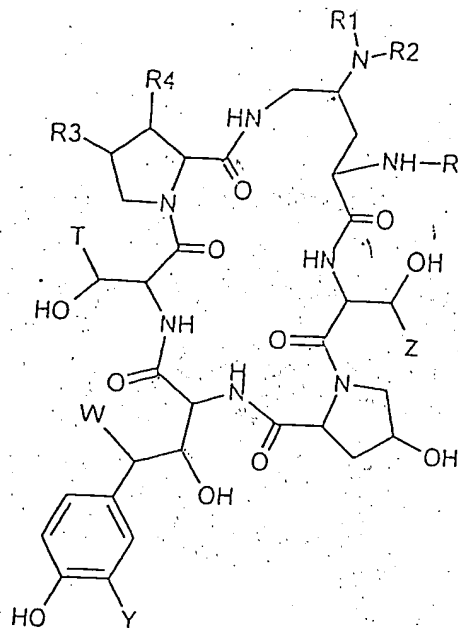


AMENDMENTS TO THE CLAIMS

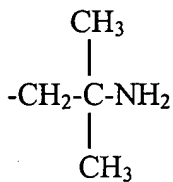
Claim 1 (currently amended)

A compound selected from the group consisting of all possible stereoisomers
~~isomeric forms and their mixtures, of~~ a compound of the formula

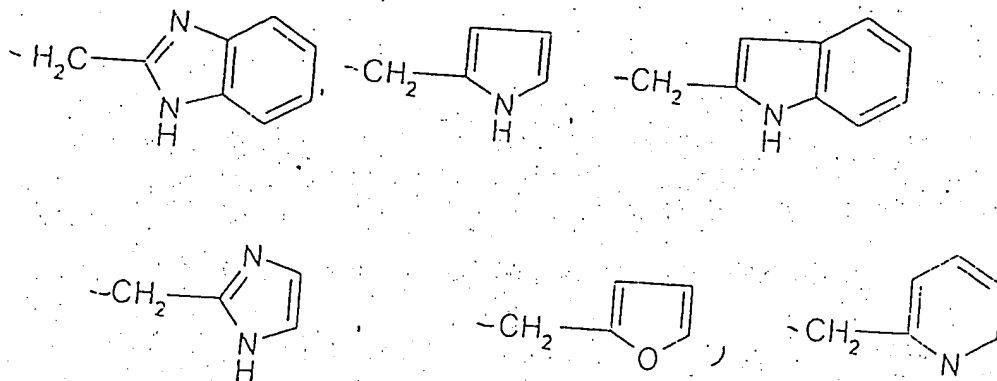


either R₁ is hydrogen or methyl and R₂ is selected from the group consisting of

-CH₂-CH₂NHCH₃,



-CH₂CHCH₃NH₂,



-CHCH₃CH₂NH₂, -(CH₂)_aOH where a is an integer of 1 to 8, -(CH₂)_b-C≡N where b is an integer of 1 to 8, -CHCH₃C₆H₅, -(CH₂)-C(CH₃)₂NHCOCF₃, and -CHCH₃(CH₂)_dOH where d is an integer of 1 to 8,

R₃ is selected from the group consisting of hydrogen, methyl and hydroxyl,

R₄ is hydrogen or hydroxyl,

R is selected from the group consisting of alkyl and cycloalkyl of up to 30 carbon atoms, optionally containing at least one heteroatom, at least one heterocycle and alkyl or cyclic acyl of up to 30 carbon atoms optionally containing at least one heteroatom, and at least one heterocycle,

T is selected from the group consisting of hydrogen, methyl, -CH₂CONH₂,

-CH₂-C≡N, and -(CH₂)₂NH₂,

Y is selected from the group consisting of hydrogen, hydroxyl, halogen and -OSO₃H or a salt thereof,

W is hydrogen or OH,

Z is hydrogen or methyl and its non-toxic, pharmaceutically acceptable acid addition salt.

Claim 2 (previously presented)

The compound of claim 1 in which T is hydrogen.

Claim 3 (previously presented)

The compound of claim 1 in which W is hydrogen.

Claim 4 (previously presented)

The compound of claim 1 in which Z is methyl.

Claim 5 (previously presented)

The compound of claim 1 in which Y is hydrogen.

Claim 6 (previously presented)

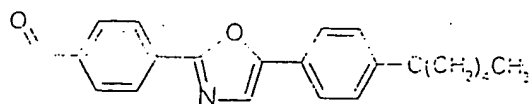
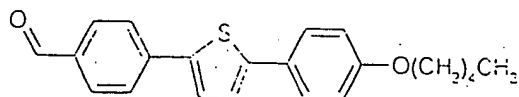
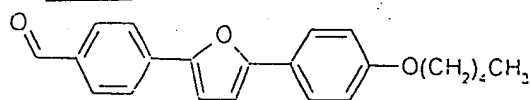
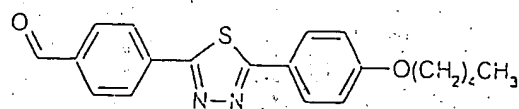
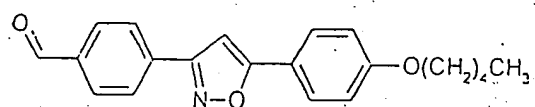
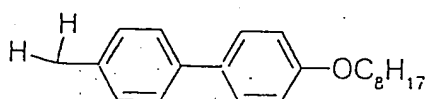
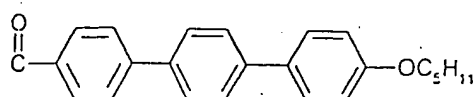
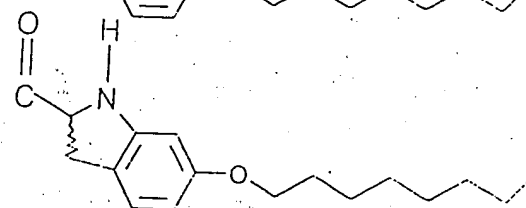
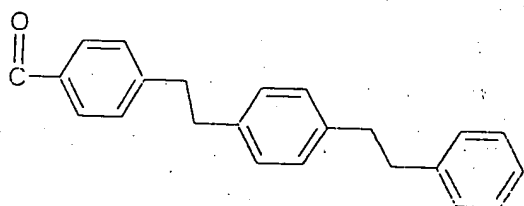
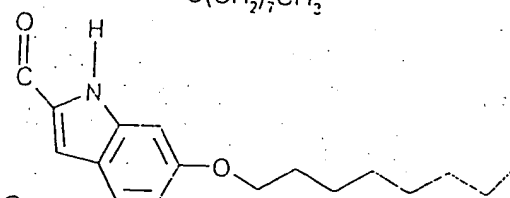
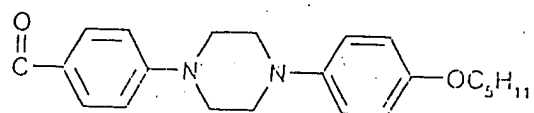
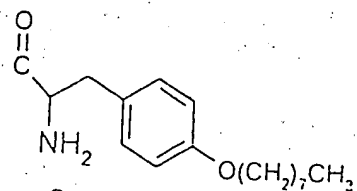
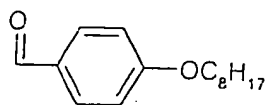
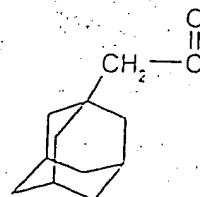
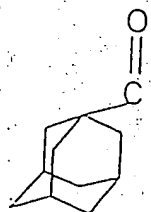
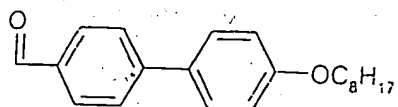
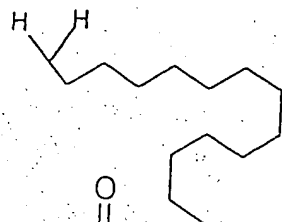
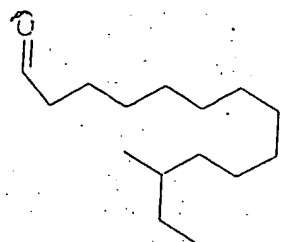
The compound of claim 1 in which R₃ is methyl.

Claim 7 (previously presented)

The compound of claim 1 in which R₄ is hydroxyl.

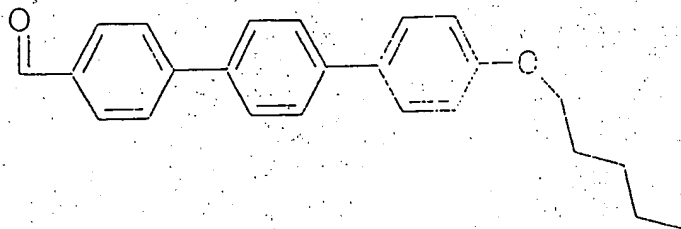
Claim 8 (previously presented)

The compound of claim 1 in which R is selected from the group consisting of



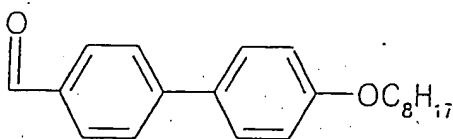
Claim 9 (previously presented)

The compound of claim 8 in which R is



Claim 10 (previously presented)

The compound of claim 8 in which R is



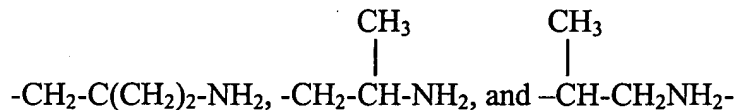
Claim 11 (previously presented)

The compound of claim 1 in which R₁ is hydrogen.

Claim 12 (cancelled)

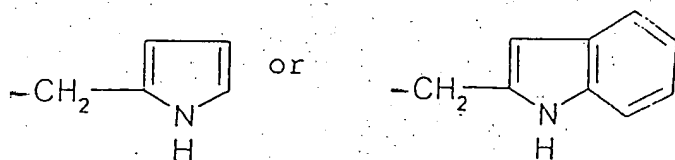
Claim 13 (previously presented)

The compound of claim 1 in which R₂ is selected from the group consisting of



Claim 14 (previously presented)

The compound of claim 1 in which R₂ is



Claim 15 (previously presented)

The compound of claim 1 is 1-[4-[[[(1H-benzimidazol-2-yl)-methyl)-amino-N2-[[4''-(pentyloxy) [1,2':4', 1''-terphenyl]-4-yl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]5-L-serine-echinocandine B trifluoroacetate (isomer B) .

Claim 16 - 18 (cancelled)

Claim 19 (previously presented)

An antifungal composition comprising an antifungally effective amount of a compound of claim 15 and an inert pharmaceutical carrier.

Claim 20 (previously presented)

A method of treating fungal infections in warm-blooded animals comprising administering to warm-blooded animals in need thereof an antifungally effective amount of a compound of claim 15.